

100
Seats Only

*CliMed Academy & Curio Training & Research Institute (CTRI), the initiatives of CliMed Research Solutions,
Bring to Ahmedabad*

MEDICAL WRITING & CLINICAL RESEARCH TRAINING WORKSHOP - AHMEDABAD 2025

August 31, 2025 (Sunday)

9.00 AM to 5.00 PM

**Training Fee Before
August 15, 2025: INR 2000/-
2000 (80% CSR-supported discount on
INR 10,000/- fee)**

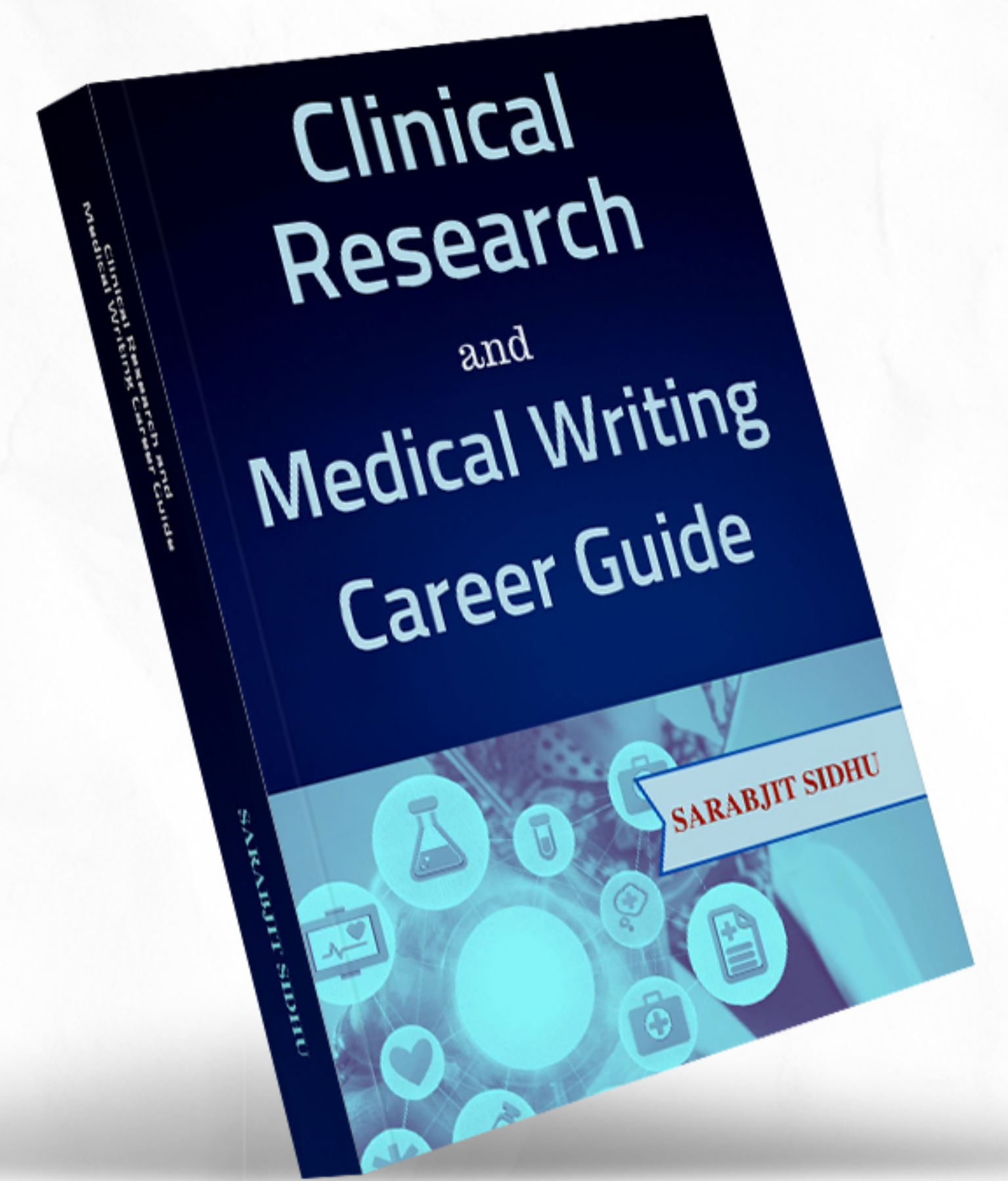


Venue:

*Hotel Platinum Inn, Ahmedabad, Near Swaminarayan Temple,
Anjali Cross Road, Vasna, Ahmedabad - 380007, Gujarat, India.*

What You'll Get:

- Free Course Material Book Hard-copy worth **599/- INR**
- Experts from the Industry to train
- One-to-One interaction with Faculty Members
- Printed Certificate of Participation
- Group Assignments & Hands-on Training
- Career Guidance & support
- High Tea & Lunch (Included)



Highlights:

- ➔ Stepwise guide to writing and publishing high-quality manuscripts
- ➔ Practical understanding of sample size, statistical tests, and data interpretation
- ➔ Insights into the future of clinical trials: Centralised and Decentralised models
- ➔ Live demonstration of core trial documents: Protocol, IB, and CSR
- ➔ Hands-on activities, real-world examples, and an interactive quiz
- ➔ Career guidance for roles in medical writing and clinical operations
- ➔ Expert-led sessions with practical takeaways for implementation

PharmD, BPharm, MPharm,
MBBS, BDS, MSc Life Sciences
students

Clinical Research
Professionals & Coordinators

Aspiring and early-career
Medical Writers

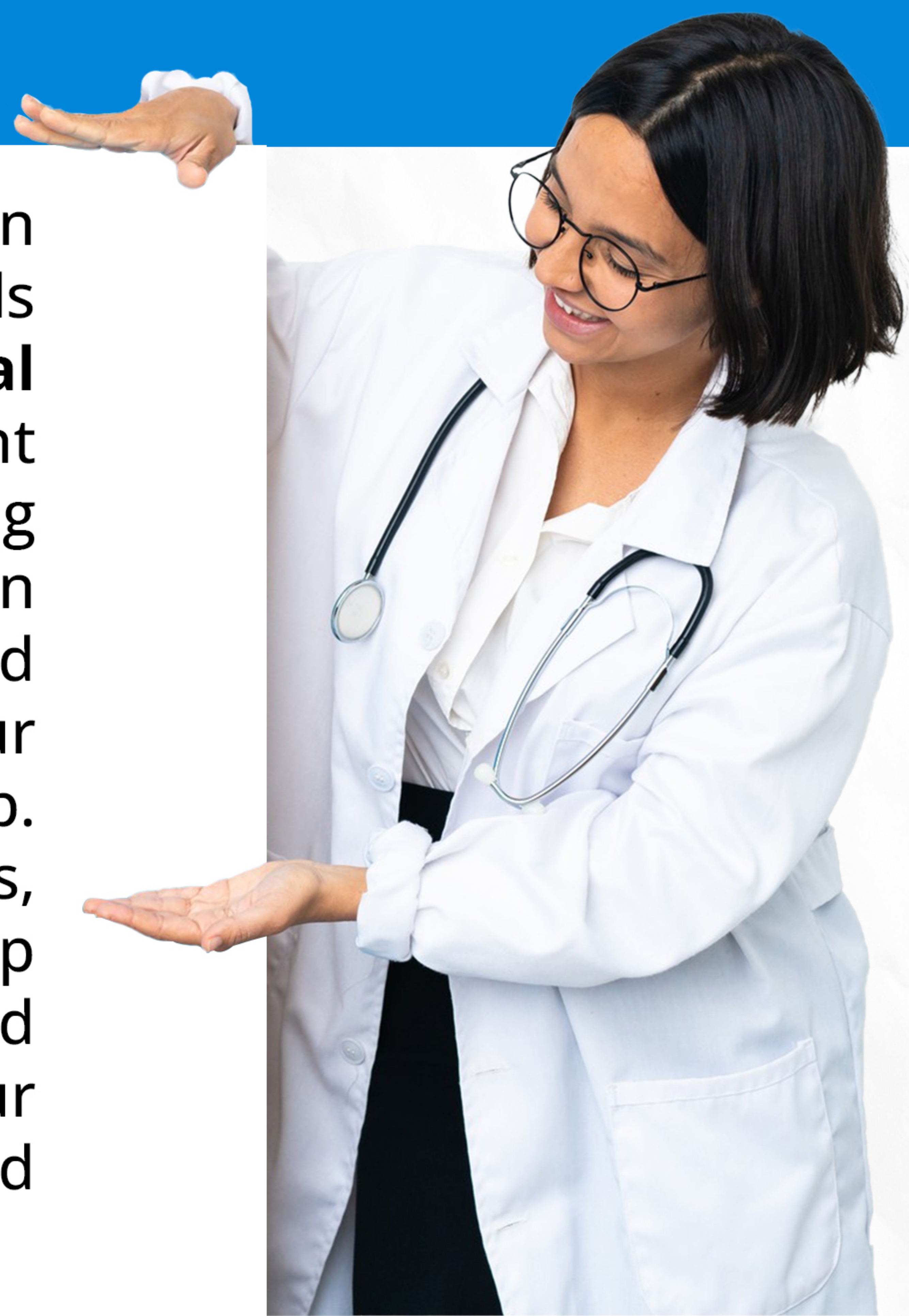
**Who
Can
Attend?**

Academic
Researchers & Postgraduates

Faculty members interested
in clinical trial education

About the Training Program:

Join us for a power-packed, hands-on workshop focused on building essential skills in **medical writing, clinical research, & trial documentation**. This one-day event features four in-depth sessions covering manuscript writing, applied statistics in clinical trials, centralised monitoring and decentralised trials (DCTs), and a 3-hour industrial medical writing workshop. Designed for early-career professionals, students, and researchers, the workshop offers real-world insights, frameworks, and interactive activities to elevate your expertise in the clinical research and publication domain.



About CliMed:

CliMed Research Solutions, founded in 2020 by Dr. Ajit Singh, is a leading Indian organization dedicated to advancing research, education, and career development in healthcare. In just five years, CliMed has trained over 100,000 students and professionals across medicine, pharmacy, nursing, and allied health sciences. Known for its innovative and practical approach, CliMed offers expertise in clinical research, pharmacovigilance, regulatory affairs, and medical writing. Through its verticals like Curio, CliMed Academy, CliMed Overseas, CARE, and CliPod, the organization provides internships, mentorships, international exam preparation, and research support. Recognized with prestigious awards like the Ratan Tata Leadership Award and Best Ed-Tech Company, CliMed continues to bridge academia and industry, fostering skill-based education and global healthcare opportunities.



Event Topics Coverage:

Session 1: How to write an acceptable manuscript for Publication?

Including

- *Types of Manuscript,*
- *Stepwise Approach,*
- *Frameworks and outlines for all types of papers, like Case reports, review papers, original papers, and systematic review papers*
- *Journal Selection & Submission*

Session 2: Suitable Statistics for Clinical Trials

Including

- *Sample Size Calculations*
- *Importance of Sample Size*
- *Descriptive Statistics*
- *Parametric and Nonparametric Tests*

Session 3: Centralised Monitoring & Decentralised Clinical Trials (DCTs)

Including

- *Introduction to Centralised Monitoring in Clinical Trials*
- *Fundamentals of Decentralised Clinical Trials (DCTs)*
- *Technological Innovations Enabling Centralised & Decentralised Trials*
- *Regulatory and Ethical Considerations in Remote Trials*
- *Operational Challenges and Best Practices for Implementation*

Three-hour Workshop

Session 4: Mastering Industrial Medical Writing with Core Clinical Trial Documents: Protocol, Investigator's Brochure, and Clinical Study Report

About Workshop: This 3-hour interactive workshop is designed to equip participants with a comprehensive understanding of three foundational documents in Medical Writing & Clinical Research as Protocol, Investigator's Brochure (IB), and Clinical Study Report (CSR). The session will include real-world examples, document structure breakdowns, and the regulatory relevance of each document. Participants will engage in hands-on activities and a quiz to reinforce learning and assess comprehension. Ideal for early-career researchers, medical writers, and clinical professionals aiming to strengthen their documentation expertise.

Workshop Coverage:

- Overview and purpose of Clinical Trial Protocol with key components
- Structure and content of the Investigator's Brochure (IB)
- Understanding Clinical Study Reports (CSR) as per ICH E3 guidelines
- Real-world examples and case-based discussion for each document
- Common challenges and quality tips in writing and reviewing these documents
- Interactive quiz to assess learning and application
- Q&A session and guidance on career roles involving these documents



Chief Guest



Prof. Pramod Kumar Rajput

*Director Board, Curio
Former Sr Vice President,
Cadila Pharma*

Chief Guest



Dr Kiran Kumar Shetty

*DGM, Meril Life Science
Pvt Ltd, Vapi, Gujarat*

Speakers



Dr. Sarabjit Sidhu

*Associate Director,
Clinical Trial
Transparency, GSK*



Mr. Kapil Dev Jhawar

*Associate Clinical
Project Management
Director, IQVIA*



Ms. Meghana Dahiya

*Manager,
Medical Writing,
Propharma*



Mr. Siddheesh Rajpurohit

*Lead Medical Writer,
Meril Life Sciences
Pvt Ltd, Vapi, Gujarat*



Dr. Ajit Singh

*Founder & CEO, CliMed & Curio
Research Scientist ICMR
Department of Medicine, Kasturba Medical College, Manipal*

Convener:



Dr. Neha Suratiya

*Deputy Director Research,
CliMed Research Solutions, India*



Day & Date:

August 31, 2025 (Sunday)
9.00 AM – 5.00 PM



Early Bird Training Fee (before August 15, 2025):

2000/- INR (80% CSR-supported
discount on INR 10000 fee)



Last Date Fee (from August 16-22, 2025):

3000/- INR (70% CSR-supported
discount on INR 10000 fee)

Early Bird Registration Link



<https://rzp.io/rzp/mwahmd>

August 31, 2025

Timings	Topic	Speaker
9.15 AM – 9.30 AM	Inauguration	Prof. Pramod Kumar Rajput
9.30 AM – 10.30 AM	How to write an acceptable manuscript for Publication?	Dr. Ajit Singh
10.30 AM – 10.45 AM	Tea Break	
10.45 AM – 11.45 AM	Suitable Statistics for Clinical Trials	Dr. Siddheesh Rajpurohit
11.45 AM – 12.45 PM	Centralised Monitoring & Decentralised Clinical Trials (DCTs)	Mr. Kapil Dev Jhawar
12.45 PM – 1.30 PM	Lunch Break	
1.30 PM – 4.30 PM	Mastering Industrial Medical Writing with Core Clinical Trial Documents: Protocol, Investigator's Brochure, and Clinical Study Report	Ms. Meghana Dahiya
		Dr. Sarabjit Sidhu
2.45 PM – 3.00 PM	High Tea	
4.30 PM	Certification Distribution & Valedictory	

